

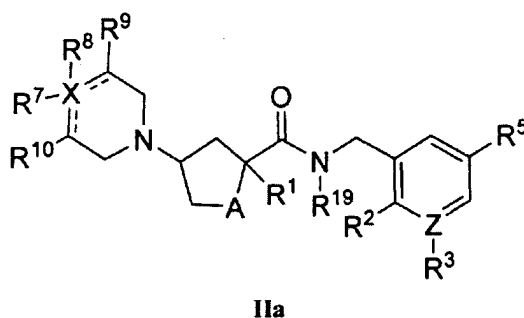
Amendments to the Claims:

This listing of Claim will replace all prior versions, and listings, of claims in the application.

Listing of claims:

Claims 1-45 (cancelled).

46. (New) A compound having the Formula IIa,



or a pharmaceutically acceptable salt thereof or an individual diastereomer thereof, wherein:

X and Z are C;

R¹ is selected from: -C₁₋₆alkyl unsubstituted or substituted with 1-6 substituents independently selected from halo, hydroxy, -O-C₁₋₃alkyl and trifluoromethyl; and phenyl unsubstituted or substituted with 1-3 substituents independently selected from halo, hydroxyl, C₁₋₃alkyl, C₁₋₃alkoxy and trifluoromethyl;

R² and R³ are independently selected from: hydrogen, fluoro, chloro, and C₁₋₃alkyl unsubstituted or substituted with 1-3 fluoro;

R⁵ is selected from fluoro, chloro, and -C₁₋₆alkyl unsubstituted or substituted with 1-6 substituents selected from fluoro;

R⁷ is fluorophenyl;

R⁸ is hydrogen;

or R⁷ and R⁸ join to form a 1H-indene ring, which is optionally substituted with 1-5 substituents independently selected from halo, trifluoromethyl, and C₁₋₃alkyl;

A is selected from -O- and -N(R²⁰)-;

R⁹ and R¹⁰ are independently selected from: hydrogen and C₁₋₆alkyl;

R¹⁹ is selected from: hydrogen and C₁₋₆ alkyl; and

R²⁰ is selected from: hydrogen, C₁₋₆ alkyl, C(=O)CH₃, and BOC.

47. (New) The compound of claim 46, wherein

R¹ is selected from: -C₁₋₆alkyl unsubstituted or substituted with 1-6 substituents independently selected from halo; and phenyl unsubstituted or substituted with 1-3 substituents independently selected from halo, C₁₋₃alkyl, and trifluoromethyl.

48. (New) The compound of claim 46, wherein R² is H.

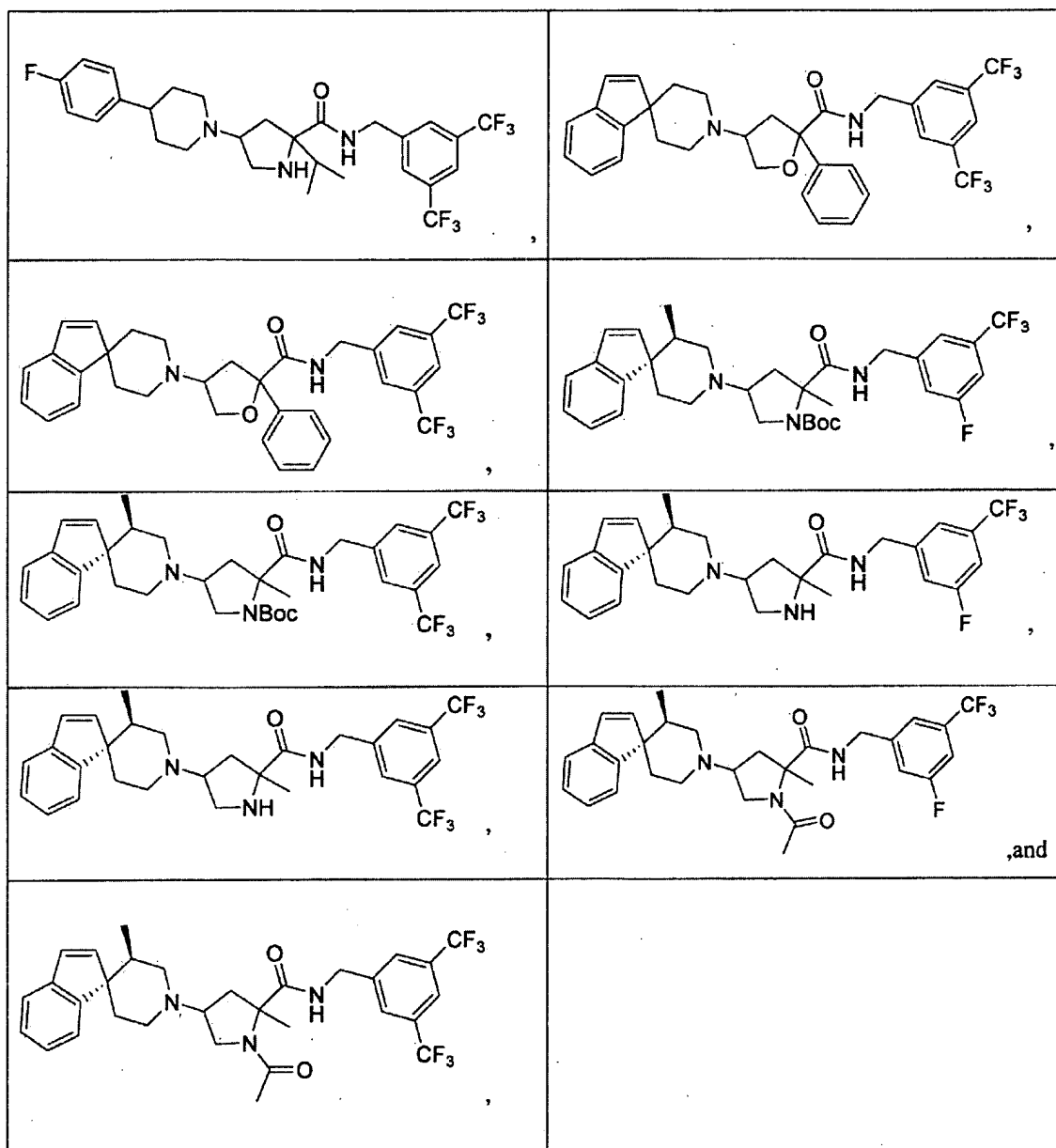
49. (New) The compound of claim 46, wherein R³ is selected from: trifluoromethyl, chloro, and fluoro.

50. (New) The compound of claim 46, wherein R⁵ is selected from hydrogen, trifluoromethyl, chloro and fluoro.

51. (New) The compound of claim 46, where R⁷ and R⁸ join to form a 1H-indene ring, wherein said ring is unsubstituted.

52. (New) The compound of Claim 46, wherein R¹⁹ and R² are H.

53. (New) A compound selected from the group consisting of:



or a pharmaceutically acceptable salt thereof, or individual diastereomer thereof.

54. (New) A pharmaceutical composition which comprises an inert carrier and the compound of Claim 46.

55. (New) A method for modulation of chemokine receptor activity in a mammal which comprises the administration of an effective amount of the compound of Claim 46.